IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

Maria Fardis et al.

Examiner:

Unknown

Serial No .:

10/583,573

Group Art Unit:

Unknown

Filed: itle:

June 19, 2006

Docket:

01692.258US2

4'-SUBSTITUTED CARBOVIR AND ABACAVIR-DERIVATIVES AS WELL AS

RELATED COMPOUNDS WITH HIV AND HCV ANTIVIRAL ACTIVITY

INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

In compliance with 37 C.F.R. § 1.56, and in accordance with 37 C.F.R. §§ 1.97 et. seq., the enclosed materials are brought to the attention of the Examiner for consideration in connection with the above-identified patent application. Applicant respectfully requests that this Information Disclosure Statement be entered and the documents listed on the attached Form 1449 be considered by the Examiner and made of record. Pursuant to MPEP 609, Applicant requests that a copy of the Form 1449, initialed as being considered by the Examiner, be returned to the Applicant with the next official communication. Applicant also encloses a copy of the corresponding International Search Report for your convenience.

Pursuant to 37 C.F.R. § 1.97, no fee or statement is required with the Information Disclosure Statement. However, the Commissioner is hereby authorized to charge the required fees to Deposit Account No. 503503 in order to have this Information Disclosure Statement considered if required. The Examiner is invited to contact the Applicant's Representative at the below-listed telephone number if there are any questions regarding this communication.

> Respectfully submitted, Maria Fardis et al. By their Representatives, Viksnins Harris & Padys PLLP P.O. Box 111098 St. Paul, MN 55111-1098 952 876-4092

Date 1/-1-06

Robert J. Harris Reg. No. 37,346

CERTIFICATION OF HAND DELIVERY: The undersigned certifies that this correspondence is being hand delivered to the United States Patent Office in an package addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450, on this day of November 2006.

(to be used for	ANSMITTAL FORM all correspondence after initial Pages in This Submission	Attorney Docket Number	01092.230	2006 rdis 8US2
Amendme Af Af Extension Express A Information Certified C Document Reply to I Incomplet	fter Final ffidavits/declaration(s) n of Time Request Abandonment Request on Disclosure Statement Copy of Priority	Drawing(s) Licensing-related Papers Petition Petition to Convert to a Provisional Application Power of Attorney, Revocat Change of Correspondence Terminal Disclaimer Request for Refund CD, Number of CD(s) Landscape Table on C	Address	After Allowance Communication to TC Appeal Communication to Board of Appeals and Interferences Appeal Communication to TC (Appeal Notice, Brief, Reply Brief) Proprietary Information Status Letter Other Enclosure(s) (please Identify below): Form 1449 (6 pgs.); Copies of 84 cited references; Return Postcard
Firm Name	SIGNA VIKSNINS HARRIS & PA	TURE OF APPLICANT, ATTO DYS PLLP	ORNEY, C	OR AGENT
Signature	100	12		
Printed name	Robert J. Harris			
Date	November, 2006 Reg. No. 37,346			
I hereby certify th sufficient postage the date shown be Signature	at this correspondence is b as first class mail in an en	ERTIFICATE OF TRANSMISS eing facsimile transmitted to the USP velope addressed to: Commissioner f	TO or depos	ILING sited with the United States Postal Service with P.O. Box 1450, Alexandria, VA 22313-1450 on

This collection of information is required by 37 CFR 1.5. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and1.14. This collection is estimated to 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Typed or printed name

Date

Substitute for form 1449A/PTO and/or 1449B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT

se as many sheets as necessary)

Complete if Known		
Application Number	10/583,573	
Filing Date	June 19, 2006	
First Named Inventor	Maria Fardis	
Group Art Unit	Unknown	
Examiner Name	Unknown	

Sheet 1 of 6

01692.258US2

US PATENT DOCUMENTS			
		Publication Date	Name of Patentee/Applicant of Document
	US 5,413,996	05-09-1995	Bodor, Nicholas S.
	US 5,670,497	09-23-1997	Bold, Guido et al.
	US 5,750,493	05-12-1998	Schinazi, Raymond F. et al.
	US 5,874,577	02-23-1999	Chen, Wei et al.
	US 5,914,332	06-22-1999	Chen, Xiaoqi et al.
	US 6,072,053	06-06-2000	Vince, Robert et al.
	US 6,312,662	11-06-2001	Robinson, Edward D. et al.
	US 5,750,343	05-12-1998	Maag, Hans et al.
	US 6,767,900	07-27-2004	Ubasawa, Masaru et al.
	US 2001-031773	10-18-2001	Camden, James Berger
	US-2003-0109498	06-12-2003	Yuasa, Satoshi et al.

FOREIGN PATENT DOCUMENTS				
Examiner Initials*	Foreign Document Number (include country code)	Publication Date	Translation (Abstract Only or Full Translation, if applicable)	
	EP 0 267 050	05-11-1988		
	EP 0 441 192	01-25-1991		
	EP 0 465 297	01-08-1992		
	EP 0 531 597	03-17-1993		
	EP 0 632 048	01-04-1995		
	EP 0 786 455	07-30-1997	1	
	EP 0 852 233	07-08-1998		
	EP 0 919 562	06-02-1999		
	EP 1 295 879	03-26-2003		
	WO 88/06158	08-25-1988		
	WO 91/19721	12-26-1991		
	WO 92/00988	01-23-1992		
	WO 92/18520	10-29-1992		
	WO 93/12123	06-24-1993		
	WO 93/24510	12-09-1993		
	WO 96/14314	05-17-1996		
	WO 96/40156	12-19-1996		
	WO 98/04569	02-05-1998		
	WO 98/11906	03-26-1998		
	WO 99/62921	12-09-1999		
	WO 00/04033	01-27-2000		
	WO 01/13957	03-01-2001		

EXAMINER

DATE CONSIDERED

Substitute for form 1449A/PTO and/or 1449B/PTO	Complete if Known		
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	Application Number	10/583,573	
NOV 0 2 2006	Filing Date	June 19, 2006	
	First Named Inventor	Maria Fardis	
	Group Art Unit	Unknown	
	Examiner Name	Unknown	
Sheet 2 of 6	01692.258US2		

PADENIA

Sheet 2 of 6

	FOREIGN PATENT DOCUMENTS			
Examiner Initials*	Foreign Document Number (include country code)	Publication Date	Translation (Abstract Only or Full Translation, if applicable)	
	WO 01/17982	03-15-2001		
	WO 01/19320	03-22-2001	,	
	WO 01/46204	06-28-2001		
	WO 01/64693	09-07-2001		
	WO 01/96329	12-20-2001		
	WO 02/03997	01-17-2002		
	WO 02/06292	01-24-2002		
	WO 02/08241	01-31-2002		
	WO 02/14344	02-21-2002		
	WO 02/057425	07-25-2002		
	WO 02/100415	12-19-2002		
	WO 03/028737	04-10-2003		
	WO 03/050129	06-19-2003		
	WO 03/059255	07-24-2003		
	WO 03/064383	08-07-2003		
	WO 03/066005	08-14-2003		
	WO 03/080078	10-02-2003		
	WO 03/090690	11-06-2003		
<u>.</u> .	WO 2004/096234	11-11-2004		
	WO 2005/011709	02-10-2005		

	OTHER DOCUMENTS NON PATENT LITERATURE DOCUMENTS
Examiner Initials*	Include last name of the first author (in CAPITAL letters), "Title of the Article", <u>Title of the Source</u> (book, magazine, journal, serial, symposium, catalog, etc.), <u>Volume-Number</u> , page(s) and (date).
muais	
	ABDEL-MEGUID, SHERIN S. et al., Inhibition of Human Immunodeficiency Virus-1 Protease
ŀ	by a C ₂ -Symmetric Phosphinate. Synthesis and Crystallographic Analysis, <i>Biochemistry</i> , 1993,
	1543-1572, Vol. 32, No. 31.
	ALLEN, LEE F. et al., CI-1040 (PDI84352), a Targeted Signal Transduction Inhibitor of MEK
	(MAPKK), Seminars in Oncology, October 2003, pp. 105-116, Vol. 30, No. 5, Elsevier Inc.
	BANTIA, SHANTA et al., Purine nucleoside phosphorylase inhibitor BCX-1777 (Immucillin-H)—
	a novel potent and orally active immunosuppressive agent, International
	Immunopharmacology, 2001, pp. 1199-1210, Elsevier Science B.V.
	BEAUCHAMP, LILIA M., et al., Guanine, Pyrazolo[3,4-d]pyrimidine, and Triazolo[4,5-
	d]pyrimidine(8-Azaguanine) Phosphonate Acyclic Derivatives as Inhibitors of Purine
	Nucleoside Phosphorylase, Journal of Medicinal Chemistry, 1996, pp. 949-956, American
	Chemical Society.
	BOHANI D. W. et al., A-420983: a potent, orally active inhibitor of lck with efficacy in a model of
	transplant rejection, Bioorganic & Medicinal Chemistry Letters, 2004, Vol. 14.

EXAMINER DATE CONSIDERED

INFORMATION DISCLOSURE	Complete if Known			
STATEMENT BY APPLICANT Use as many sheets as necessary) NOV 0 2 2006 Sheet 3 of 6	l Application Number	10/583,573		
	Filing Date	June 19, 2006		
	First Named Inventor	Maria Fardis		
	Group Art Unit	Unknown		
	Examiner Name	Unknown		
	01692.258US2			

	OTHER DOCUMENTS NON PATENT LITERATURE DOCUMENTS
Examiner Initials*	Include last name of the first author (in CAPITAL letters), "Title of the Article", <u>Title of the Source</u> (book, magazine, journal, serial, symposium, catalog, etc.), <u>Volume-Number</u> , page(s) and (date).
	BZOWSKA, AGNIESZKA et al., Purine nucleoside phosphorylases: properties, functions, and clinical aspects, <i>Pharmacology & Therapeutics</i> , 2000, pp. 349-425, Vol. 88, Elsevier Science
	Inc.
	CHAPMAN, H. et al., Practical Synthesis, Separation, and Stereochemical Assignment of the PMPA Pro-Drug GS-7340, Nucleosides, Nucleotides & Nucleic Acids, 2001, pp. 621-628, Vol. 20, Nos. 4-7, Marcel Dekker, Inc.
	CLARK, JEREMY L. et al.,
	Mycophenolic Acid Analogues as Potential Agents Against West Nile Virus Infection.
	CONKLYN, MARYROSE et al., The JAK3 inhibitor CP-690550 selectively reduces NK and
	CD8+ cell numbers in cynomolgus monkey blood following chronic oral dosing, <i>Journal of Leukocyte Biology</i> , December 2004, pp. 1-8, Vol. 76, The Society for Leukocyte Biology.
	DE CLEREQ, E., Highlights in the Development of New Antiviral Agents, <i>Mini Reviews in Medicinal Chemistry</i> , 2002, 163-175, Vol. 2, No. 2., Bentham Science Publishers, Ltd.
	DE CLERCQ, ERIK, New Developments in Anti-HIV Chemotherapy, Current Medicinal Chemistry, 2001, 1543-1572, Vol. 8, No. 13, Bentham Science Publishers Ltd.
	DVORAKOVA, HANA <i>et al.</i> , Synthesis of 2'-Aminomethyl Derivatives of N-(2-(Phosphonomethoxy)ethyl) Nucleotide Analogues as Potential Antiviral Agents, <i>J. Med. Chem.</i> , 1996, 3263-3268. Vol. 38, No. 17.
	EVANS, GARY B., Exploring Structure Activity Relationships of Transition State Analogues of Human Purine Nucleoside Phosphorylase, <i>J. Med. Chem.</i> , 2003, 3412-3423, Vol. 46, No. 15, American Chemical Society.
-	GUMINA, GIUSEPPE et al., Advances in antiviral agents for hepatitis B virus, Antiviral Chemistry & Chemotherapy, 2001, 93-112, Vol. 12, Suppl. 1, International Medical Press.
	GOBEC, S. et al., Phosphonate inhibitors of antiget 85C, a crucial enzyme involved in the biosynthesis of the mycobacterium tuberculosis cell wall, <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, Vol. 14.
	HEGEDUS, LOUIS S. et al., Synthesis of 4'-Methyl and 4'-cyano Carbocyclic 2',3'-Didehydro Nucleoside Analogues via 1,4-Addition to Substituted Cyclopentenones, <i>J. Org. Chem.</i> , 2004, 8492-8495, Vol. 69, No. 24, American Chemical Society.
	HERCZEGH P., et al., Osteoadsorptive bisphosphonate derivatives of fluoroquinolone antibacterials, J. Med. Chem., 2002, Vol. 45.
	HIRABAYASHI, HIDEKI et al., Bone-Specific Drug Delivery Systems, Clinical Pharacokinetics, 2003, 1319-1330, Vol. 42, No. 15.
	HOLY A. et al., Synthesis, Cliect. Czech. Chem. Commun., 1989, Vol. 54, pages 2190-2210.
	JAIN, JUGNU et al., Characterization of Pharmacological Efficacy of VX-148, a New, Potent Immunosuppressive Inosine 5'-Monophosphate Dehydrogenase Inhibitor, Journal of Pharmacology and Experimental Therapeutics, 2002, 1272-1277, Vol. 302, No. 3, The American Society for Pharmacology and Experimental Therapeutics.
	KARPENKO, INNA L. et al., Synthesis and Antitherpetic Activity of Acyclovir Phosphonates, NUCLEOSIDES, NUCLEOTIDES & NUCLEIC ACIDS, 2003, 319-328, Vol 22, No. 3, Marcel Dekker, Inc.

DATE CONSIDERED

Substitute for form 1449A/PTO and/or 1449B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT OR STATEMENT BY APPLICANT BY	Complete if Known		
	Application Number	10/583,573	
	Filing Date	June 19, 2006	
	First Named Inventor	Maria Fardis	
	Group Art Unit	Unknown	
	Examiner Name	Unknown	
	01692.258US2		

	OTHER DOCUMENTS NON DATENT LITERATURE DOCUMENTS
	OTHER DOCUMENTS NON PATENT LITERATURE DOCUMENTS Include last name of the first author (in CAPITAL letters), "Title of the Article", Title of the Source
Examiner Initials*	(book, magazine, journal, serial, symposium, catalog, etc.), Volume-Number, page(s) and (date).
	KATO, KEISUKE et al., Stereoselective synthesis of 4' - alphaalkyclcarbovir derivatives
	based on an asymmetric synthesis or chemo-enzymatic procedure, Chemical &
	Pharmaceutical Bulletin, 1999, 1256-1264, Vol. 49, No. 9, Pharmaceutical Society of Japan.
	KATO, KEISUKE et al., Enantio- and diastereoselective syntheis of 4'-α-substituted carbocyclic
	nucleosides, Tetrahedron: Asymmetry, 1998, 911-914, Vol. 9, Elsevier Science Ltd.
	KILPATRICK, J. MICHAEL, Intravenous and oral pharmacokinetic study of BCX-1777, a novel
	purine nucleoside phosphorylase transition-state inhibitor, In vivo effects on blood 2'-
	deoxyguanosine in primates, <i>International Immunopharmacology</i> , 2003, 541-548, Vol. 3,
	Elsevier Science B.V.
	KIM, CHOUNG UN et al., Regiospecific and Highly Stereoselective Electrophilic Addition to
	Furanoid Glycals: Synthesis of Phosphonate Nucleotide Analogues with Potent Activity against
	HIV, J. Org. Chem., 1991, 2642-2647, Vol. 56, No. 8, American Chemical Society.
	KINSKY, STEPHEN C. et al., Inhibition of cell proliferation by putative metabolites and non-
	degradable analogs of methotrexategamadimyristoylphosphatidylethanolamine, <i>Biochimica</i>
	et Biphysica Acta, 19878, 211-218, Vol. 917, No. 2., Elsevier Science Publishers B. V.
	KINSKY, STEPHEN C. et al., Effect of liposomes sentitized with methotrexate-γ-
	dimyristoylphosphatidylethanolamine on cells that are resistant to methotrexate, <i>Biochimica et Biophysica Acta</i> , 1986, 129-135, Vol. 885, Elsevier Science Publishers B.V.
	KINSKY, STEPHEN C. <i>et al.</i> , Circumvention of the methotrexate transport system by
	methotrexate-phosphatidylethanolamine derivatives effect of fatty acid chain length, <i>Biochimica</i>
	et Biophysica Acta, 1987, 96-103, Vol. 921, Elsevier Science Publishers B.V.
	KO, OK HYUN <i>et al.</i> , Efficient synthesis of novel carbocyclic nucleosides via sequential
	Claisen rearrangement and ring-closing metathesis, <i>Tetrahedron Letters</i> , 2002, 6399-6402,
	Vol. 43, Elsevier Science Ltd.
	LEWANDOWICZ, ANDRZEJ et al., Achieving the Ultimate Physiological Goal in Transition
	State Analogue Inhibitors for Purine Nucleoside Phosphorylase, <i>The Journal of Biological</i>
	Chemistry, 2003, 31465-31468, Vol. 278, No. 34, The American Society for Biochemistry and
1	Molecular Biology, Inc.
	MENENDEZ-ARIAS, LUIS et al. Targeting HIV: antiretroviral therapy and development of drug
	resistance, TRENDS in Pharmacological Sciences, 2002, 381-388, Vol. 23, No. 8, Elsevier
	Science Ltd.
	ONO-NITA, SUZANE KIOKO et al., Novel Nucleoside Analogue MCC-478 (LY582563) Is
	Effective against Wild-Type or Lamivudine-Resistant Hepatitis B Virus, Antimicrobial Agents
	and Chemotherapy, 2002, 2602-2605, Vol. 46, No. 8, American Society for Microbiology.
	PANKIEWICZ, KRZYSZTOF W., Novel Mycophenolic Adenine Bis(phosphonate) Analogues
	As Potential Differentiation Agents against Human Leukemia, J. Med. Chem., 2002 703-712,
	Vol. 45, No. 3, American Chemical Society.
	PARANG, KEYKAVOUS et al., Novel Approaches for Designing 5'-O-Ester Prodrugs of 3'-
	Azido-2', 3'-dideoxythymidine (AZT), Current Medicinal Chemistry, 2000, 995-1039, Vol. 7, No.
	10, Bentham Science Publishers Ltd.
	PRASHAD, MAHAVIR et al., An Efficient and Large-Scale Enantioselective Synthesis of
	PNP405: A Purine Nucleoside Phosphorylase Inhibitor, <i>J. Org. Chem.</i> , 2002, 6612-6617, Vol.
L	67, No. 19, American Chemical Society.

EXAMINER DATE CONSIDERED

Substitute for form 1449A/PTO and/or 1449B/PTO	Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	Application Number	10/583,573
STATEMENT BY APPLICANT See as many sheets as necessary) NOV 0 2 2006 Sheet 5 of 6	Filing Date	June 19, 2006
	First Named Inventor	Maria Fardis
	Group Art Unit	Unknown
	Examiner Name	Unknown
	01692.258US2	

	OTHER DOCUMENTS NON PATENT LITERATURE DOCUMENTS	
Examiner Initials*	Include last name of the first author (in CAPITAL letters), "Title of the Article", <u>Title of the Source</u> (book, magazine, journal, serial, symposium, catalog, etc.), <u>Volume-Number</u> , page(s) and (date).	
	RAY, ADRIAN S. et al., Role of Purine Nucleoside Phosphorylase in Interactions between 2',	
	3'-Dideoxyinosine and Allopurinal, Ganciclovir, or Tenofovir, Antimicrobial Agents and	
	Chemotherapy, 2004, 1089-1095, Vol. 48, No. 4, American Society for Microbiology.	
	REED, LEFF et al., Antidiabetic PPARy Ligands: An update on Compounds in development,	
	Curr. Med. Chem Imun., Endoc. & Metab. Agents, 2002, 33-47, Vol. 2, No. 1, Bentham	
	Science Publishers Ltd.	
	ROBERTS, STANLEY M., Development of the route to the new anti-AIDS drug abacavir: A highlight of academic/industry laison, <i>IDrugs</i> , 1998, 896-899, Vol. 1, No. 8, Current Drugs Ltd.	
	ROSOWSKY, ANDRE et al., METHOTREXATE ANALOGUES—27, Biochemical	
	Pharmacology, 1986, 3327-3333, Vol. 35, No. 19, Pergamon Journals Ltd.	
	ROSOWSKY, ANDRE et al., Methotrexate Analogues, 32, Chain Extension, α-Carboxyl	
	Replacement by Sulfonate and Phosphonate: Effect on Enzyme Binding and Cell-Growth	
	Inhibition, J. Med. Chem., 1988, 1326-1331, Vol. 31, No. 7, American Chemical Society.	
	SCHULTZ, C., Prodrugs of biologically active phosphate esters, Bioorganic & Medicinal	
	Chemistry, 2003, 885-898, Vol. 11, Elsevier Science Ltd., GB.	
	SEKIYA, KOUICHI et al., 2-Amino-6-arylthio-9-[2-(phosphonomethoxy) ethyl) purine Bis(2,2,2-	
	trifluoroethyl) Esters as Novel HBV-Specific Antiviral Reagents, Journal of Medicinal	
	Chemistry, 2002, 3138-3142, Vol. 45, No. 14, American Chemical Society.	
	SHI, WUXIAN et al., Plasmodium falciparum Purine Nucleoside Phosphorylase, The Journal of	
	Biological Chemistry, 2004, 18103-18106, Vol. 279, No. 18, The American Society of	
	Biochemistry and Molecular Biology, Inc.	
	SINTCHAK, MICHAEL D. et al., The structure of inosine 5'-monophosphate dehydrogenase	
	and the design of novel inhibitors, Immunopharmachology, 2000, 163-184, Vol. 47, Elsevier.	
	SRINIVAS, RANGA V. et al., Metabolism and In Vitro Antiretroviral Activities of	
	Bis(Pivaloyloxymethyl) Prodrugs of Acyclic Nucleoside Phosphonates, Antimicrobial Agents	
	and Chemotherapy, 1993, 2247-2250, Vol. 37, No. 10, American Society for Microbiology.	
	STURTZ, GEORGES <i>et al.</i> , Su rune nouvelle approche de pharmacomodulation du methotrexate: synthese d'analogues gem-diphosphoniques d'amethopterine et de la N-10	
	deaza amethopterine, Medicinal Chemistry, C. R. Acad. Sci. Paris, 1990, Vol. 10, No. 2, 739-	
	742, Academie des Sciences.	
	STURTZ, GEORGES <i>et al.</i> , Analogues phosphonoglutamiques d'amethopterine	
	(methotrexate), Eur. J. Med. Chem – Chim. Ther., 1984, 267-273, Vol. 19, No. 3.	
	STURTZ, G. et al., Synthesis of gem-bisphosphonic methotrexate conjugates and their	
	biological response towards Walker's osteosarcoma, <i>Eur. J. Med. Chem.</i> , 1993, 899-903, Vol.	
	28, Elsevier.	
	STURTZ, G. et al., A study of the delivery-targeting concept applied to antineoplasic drugs	
	active on human osteosarcoma, I. Synthesis and biological activity in nude mice carrying	
	human osteosarcoma xenografts of gem-bisphosphonic methotrexate analogues, Eur J. Med.	
	Chem., 1992, 825-833, Vol. 27, No. 8, Elsevier.	
	VIELHABER, BERND, Bericht vom 3rd International Workshop on Salvage Therapy for HIV-	
	Infection, Deutsche Aids-Hilfe e.V. FaxReport zu HIV und AIDS, 2000, 12-14.	
	WAEGELL W. et al. A420983, a novel, small molecule inhibitor of LCK prevents allograft	
	rejection, Transplantation Proceedings, 2002, 1411-1417, Vol. 34.	

EXAMINER DATE CONSIDERED

Substitute for form 1449A/PTO and/or 1449B/PTO Complete if Known INFORMATION DISCLOSURE 10/583,573 **Application Number** ATATEMENT BY APPLICANT (USA) at many sheets as necessary) June 19, 2006 **Filing Date First Named Inventor** Maria Fardis **Group Art Unit** Unknown Unknown **Examiner Name** 01692.258US2 Sheet 6 of 6

	OTHER DOCUMENTS NON PATENT LITERATURE DOCUMENTS
Examiner Initials*	Include last name of the first author (in CAPITAL letters), "Title of the Article", Title of the Source (book, magazine, journal, serial, symposium, catalog, etc.), Volume-Number, page(s) and (date).
	WROBLEWSKI, ANDRZEJ <i>et al.</i> , Synthesis of (1R,2S)- and (1S,2S)-3-(4-carbamoyl-1,2,3-triazol-1-yl)-1,2-dihydroxypropylphosphonates, Tetrahedron: Asymmetry, 2004, 1457-1464,
	Vol. 15, Elsevier.